I CLAIM:

1. A compound of the formula

$$R^4$$
— $(CH_2)_n$ — Y
 R^0
 R^3
 R^1
 (I)

wherein

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 R^1 is -H, -OH, -O(C1-C4 alkyl), -OCOC6H5, -OCO(C1-C6 alkyl), or -OSO2(C2-C6 alkyl);

10 R^0 , R^2 and R^3 are each independently -H, -OH, -O(C₁-C₄ alkyl), -OCOC₆H₅, -OCO(C₁-C₆ alkyl), -OSO₂(C₂-C₆ alkyl) or halo;

R⁴ is 1-piperidinyl, 1-pyrrolidinyl, methyl-1-pyrrolidinyl, dimethyl-1-pyrrolidinyl, 4-morpholino, dimethylamino, diethylamino, diisopropylamino, or 1-hexamethyleneimino;

15 n is 2 or 3;

X is -S- or -HC=CH-; and

Y is -O-, -S-, -NH-, -NMe-, or -CH₂-;

or a pharmaceutically acceptable salt thereof.

2. A compound of Claim 1 of the formula

$$R^4$$
— $(CH_2)_n$ — Y
 R^3

wherein

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5 R^1 is -H, -OH, -O(C₁-C₄ alkyl), -OCOC₆H₅, -OCO(C₁-C₆ alkyl), or -OSO₂(C₂-C₆ alkyl);

 R^2 and R^3 are each independently -H, -OH, -O(C₁-C₄ alkyl), -OCOC₆H₅, -OCO(C₁-C₆ alkyl), -OSO₂(C₂-C₆ alkyl) or halo;

R⁴ is 1-piperidinyl, 1-pyrrolidinyl, methyl-1-pyrrolidinyl, dimethyl-1-pyrrolidinyl, 4-morpholino, dimethylamino, diethylamino, diisopropylamino, or 1-hexamethyleneimino;

n is 2 or 3;

X is -S- or -HC=CH-; and

Y is -O-, -S-, -NH-, -NMe-, or -CH₂-;

- or a pharmaceutically acceptable salt thereof.
 - 3. A compound according to Claims 1 or 2 wherein Y is -O-.
 - 4. A compound according to any of Claims 1 to 3 wherein n is 2.

5. A compound according to any of Claims 1 to 4 wherein R¹ is -OH or -OCH₃.

- 6. A compound according to any of Claims 1 to 5 wherein \mathbb{R}^1 is -OH.
- 7. A compound according to any of Claims 1 to 6 wherein R⁴ is 1-piperidinyl or 1-pyrrolidinyl.

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- 8. A compound according to any of Claims 1 to 7 wherein R⁴ is 1-piperidinyl.
- 9. A compound according to any of Claims 1 to 8 wherein two of R⁰, R² and 5 R³ are -H.
 - 10. A compound according to any of Claims 1 to 8 wherein two of R^0 , R^2 and R^3 are -H and the other is -OH.
- 10 11. A compound according to any of Claims 1 to 9 wherein R⁰, R² and R³ are all -H.
 - 12. A compound according to any of Claims 1 to 11 wherein X is -S-.
- 15 13. A compound according to any of Claims 1 to 12 wherein X is -HC=CH-.
 - 14. A compound according to Claim 1 wherein said compound is selected from the group consisting of:
- 11-[4-(2-Piperidin-1-yl-ethoxy)-phenyl]-6,11-dihydro-benzo[b]naphtho[2,3-20 d]thiophen-3-ol;
 - 11-[4-(2-Piperidin-1-yl-ethoxy)-phenyl]-6,11-dihydro-benzo[b]naphtho[2,3-d]thiophene-3,8-diol;
 - 11-[4-(2-Piperidin-1-yl-ethoxy)-phenyl]-6,11-dihydro-benzo[b]naphtho[2,3-d]thiophene-3,10-diol;
- 25 11-[4-(2-Piperidin-1-yl-ethoxy)-phenyl]-6,11-dihydro-benzo[b]naphtho[2,3-d]thiophene-3,9-diol;
 - 10-Fluoro-12-[4-(2-piperidin-1-yl-ethoxy)-phenyl]-7,12-dihydrobenzo[a]anthracen-3-ol;
- 12-[4-(2-Piperidin-1-yl-ethoxy)-phenyl]-7,12-dihydro-benzo[a]anthracene-3,10-30 diol;
 - 12-[4-(2-Piperidin-1-yl-ethoxy)-phenyl]-7,12-dihydro-benzo[a]anthracene-3,9-diol;

- 12-[4-(2-Piperidin-1-yl-ethoxy)-phenyl]-7,12-dihydro-benzo[a]anthracene-3,11-diol;
- 12-[4-(2-Piperidin-1-yl-ethoxy)-phenyl]-7,12-dihydro-benzo[a]anthracene-3,8-diol;
- 5 or a pharmaceutically acceptable salt thereof.
 - 15. A compound according to Claim 1 wherein said compound is 12-[4-(2-Piperidin-1-yl-ethoxy)-phenyl]-7,12-dihydro-benzo[a]anthracene-3,9-diol or a pharmaceutically acceptable salt thereof.

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16. A pharmaceutical composition comprising a compound according to any of Claims 1 to 15 or a pharmaceutically acceptable salt thereof, and optionally an effective amount of estrogen and progestin, in combination with a pharmaceutically acceptable salt, diluent, or excipient.

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- 17. A method for inhibiting a disease associated with estrogen deprivation comprising administering to a patient in need thereof a therapeutically effective amount of a compound according to any one of Claims 1 through 15.
- 20 18. A method according to Claim 17 wherein said patient is a human.
 - 19. A method according to Claim 18 wherein said patient is a postmenopausal female.
- 25 20. A method according to any of Claims 17 to 19 wherein said disease is bone loss.
 - 21. A method according to any of Claims 17 through 19 wherein said disease is cardiovascular disease.

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22. A method for inhibiting a disease associated with an aberrant physiological response to endogenous estrogen comprising administering to a patient in need thereof a

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therapeutically effective amount of a compound according to any one of Claims 1 through 15.

- 23. A method according to Claim 22 wherein said patient is a human.
- 24. A method according to Claim 23 wherein said patient is a postmenopausal female.
- 25. A method according to any of Claims 22 through 24 wherein the disease associated with an aberrant physiological response to endogenous estrogen is estrogen dependent cancer.
 - 26. A method according to Claim 25 wherein said cancer is breast cancer.
- 27. A method according to any of Claims 22 through 24 wherein the disease associated with an aberrant physiological response to endogenous estrogen is endometriosis.
- 28. A method according to any of Claims 22 through 24 wherein the disease associated with an aberrant physiological response to endogenous estrogen is uterine fibrosis.
 - 29. The use of a compound according to any of Claims 1 to 15 for the manufacture of a medicament.
 - 30. The use of a compound according to any of Claims 1 to 15 for the manufacture of a medicament for inhibiting a disease associated with estrogen deprivation.
- 30 31. The use according to Claim 30 wherein said disease is bone loss.

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- 32. The use according to Claim 30 wherein said disease is cardiovascular disease.
- 33. The use of a compound according to any of Claims 1 to 15 for the manufacture of a medicament for inhibiting a disease associated with an aberrant physiological response to endogenous estrogen.
- 34. The use according to Claim 33 wherein said disease is estrogen dependent cancer.
- 10 35. The use according to Claim 34 wherein said cancer is breast cancer.
 - 36. The use according to Claim 33 wherein the disease associated with an aberrant physiological response to endogenous estrogen is endometriosis.
- 15 37. The use according to Claim 33 wherein the disease associated with aberrant physiological response to endogenous estrogen is uterine fibrosis.
 - 38. A pharmaceutical composition for inhibiting a disease associated with deprivation containing as an active ingredient a compound according to Claims 1 to 15.
 - 39. A pharmaceutical composition for inhibiting a disease associated with an aberrant physiological response to endogenous estrogen containing as an active ingredient a compound according to any of Claims 1 to 15.

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